

Department of Vermont Health Access Pharmacy Benefits Management Program DUR Board Meeting Agenda

September 10, 2019: 6:30 – 8:30 p.m.

| September 10, 2017. 0.30 – 0.30 p.m. | | |
|--------------------------------------|---|-------------|
| • | Executive Session | 6:00 - 6:30 |
| • | Introductions and Approval of DUR Board Minutes (Public Comment Prior to Board Action) | 6:30 - 6:35 |
| • | DVHA Pharmacy Administration Updates | 6:35 - 6:40 |
| • | Medical Director Update | 6:40 - 6:45 |
| • | Follow-up Items from Previous Meetings | 6:45 - 6:45 |
| • | RetroDUR/ProDUR Introduce: Discussion Topics for 2020 RetroDUR Initiatives Data Presentation: Use of Gabapentin | 6:45 - 7:15 |
| • | Clinical Update: Drug Reviews (Public comment prior to Board action) | 7:15-7:45 |

Biosimilar Drug Reviews

None at this time

Full New Drug Reviews

(Any new drug reviews that also fall within the Therapeutic Class review will be discussed during the Therapeutic Class Review)

- Apadaz® (benzhydrocodone and acetaminophen)
- Firdapse® (amifampridine)
- Gamifant® (emapalumab-lzsg)
- Inbrija® (levodopa inhalation powder)
- Lexette® (halobetasol propionate)
- Motegrity® (prucalopride)
- Nuzyra® (omadacycline)
- Qbrexza® (glycopyrronium)
- Seysara® (sarecycline tablets)
- Spravato® (esketamine nasal spray)
- Symjepi® (epinephrine)

New Managed Therapeutic Drug Classes

(Public comment prior to Board action)

None at this time

7:45 - 7:45

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| Antiemetics | | |
| Antipsychotics | | |
| Bronchodilators: Beta Agonists | | |
| COPD Agents | | |
| Growth Hormones | | |
| Inhaled Glucocorticosteroids | | |
| PAH | | |
| Review of Newly-Developed/Revised Criteria (Public comment prior to Board action) None at this time | 8:25 - 8:25 | |
| General Announcements | 8:25-8:30 | |
| Selected FDA Safety Alerts | | |
| FDA approves Boxed Warning about increased risk of blood clots and | | |
| higher dose of arthritis and ulcerative colitis medicine tofacitinib (Xeljanz, | | |
| Xeljanz XR) | , , , , , , , , , , , , , , , , , , , | |
| https://www.fda.gov/drugs/drug-safety-ar | nd-availability/fda-approves-boxed- | |
| warning-about-increased-risk-blood-clots | | |
| warming-about-mereased-msk-blood-clots | -and-deadi-inglier-dose-artificity-and | |

7:45 - 8:25

8:30

Therapeutic Drug Classes – Periodic Review (Public comment prior to Board action)

Androgenic Agents

Adjourn